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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/850,293	05/07/2001	Robert Falotico	CRD-0931	2210	
27777	7590 10/16/2003		EXAMINER		
PHILIP S. JOHNSON			ODLAND, KA	ODLAND, KATHRYN P	
JOHNSON & JOHNSON ONE JOHNSON & JOHNSON PLAZA NEW BRUNSWICK, NJ 08933-7003		ART UNIT	PAPER NUMBER		
			3743	4.00 /	
			DATE MAILED: 10/16/2003		

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)			
Office Action Summary		09/850,293	FALOTICO, ROBERT			
		Examiner	Art Unit			
		Kathryn Odland	3743			
	DATE of this communication ap	opears on the cover sheet with the				
Period for Reply						
THE MAILING DATE - Extensions of time may be a after SIX (6) MONTHS from - If the period for reply specif - If NO period for reply is specif - Failure to reply within the second reply received by the Operiod patent term adjustment	OF THIS COMMUNICATION available under the provisions of 37 CFR 1 the mailing date of this communication. ied above is less than thirty (30) days, a recified above, the maximum statutory period of or extended period for reply will, by statu ffice later than three months after the mailing	LY IS SET TO EXPIRE 3 MONT	timely filed days will be considered timely. om the mailing date of this communication. NED (35 U.S.C. § 133).			
Status 1) Responsive to	o communication(s) filed on 20	August 2002				
	communication(s) filed on 29	-				
2a) ☐ This action is	·	This action is non-final.				
		vance except for formal matters, or <i>Ex parte Quayle</i> , 1935 C.D. 11	•			
·	is/are pending in the application	on.				
	4a) Of the above claim(s) <u>14</u> is/are withdrawn from consideration.					
	Claim(s) is/are allowed.					
	⊠ Claim(s) <u>1-14</u> is/are rejected.					
	Claim(s) is/are objected to.					
8) Claim(s)	are subject to restriction and	or election requirement.				
Application Papers						
9) The specificatio	n is objected to by the Examin	ier.				
10) The drawing(s)	filed on is/are: a)□ acc	epted or b) objected to by the Ex	xaminer.			
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
11) The proposed drawing correction filed on is: a) approved b) disapproved by the Examiner.						
If approved, corrected drawings are required in reply to this Office action.						
12) The oath or declaration is objected to by the Examiner.						
Priority under 35 U.S.C.	. §§ 119 and 120					
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a)□ All b)□ So —	me * c) None of:					
1.☐ Certified	1. Certified copies of the priority documents have been received.					
2.☐ Certified	2. Certified copies of the priority documents have been received in Application No					
appli	cation from the International B	ority documents have been rece Sureau (PCT Rule 17.2(a)). St of the certified copies not rece	•			
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).						
a) ☐ The translation of the foreign language provisional application has been received. 15) ☑ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.						
Attachment(s)		p				
1) Notice of References Cit 2) Notice of Draftsperson's	ed (PTO-892) Patent Drawing Review (PTO-948) tatement(s) (PTO-1449) Paper No(s)	5) Notice of Inform	ary (PTO-413) Paper No(s) al Patent Application (PTO-152)			

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DETAILED ACTION

Response to Amendment

This is a response to the amendment dated August 29, 2003. Claims 1-14 are pending. Claim 15 has been cancelled.

Response to Arguments

1. Applicant's arguments with respect to claims 1 and 8 have been considered but are most in view of the new ground(s) of rejection. Applicant's amendments have changed the scope of the claim; thus, a new rejection has been applied.

Claim Rejections - 35 USC § 103

- 2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 3. Claims 1-14 are rejected under 35 U.S.C. 103(a) as being unpatentable over Chudzik et al. in US Patent No. 6,214,901 in view of Morris et al. in US Patent No. 5,516,781.

Regarding claims 1 and 8, Chudzik et al. disclose a method/drug delivery device for preventing/treating constrictive vascular remodeling via a controlled delivery, by release from a stent, of a compound in therapeutic dosage amounts in the range from about thirty-five micrograms per fifteen millimeters of stent to about four hundred thirty micrograms per fifteen to eighteen millimeters of stent, the compound substantially reducing in-lesion lumen loss both proximate and distal

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to the stent, the compound being incorporated in a polymeric matrix, as recited in column 1, lines 20-32, lines 42-46, lines 55-60, column 3, lines 1-46, column 5, lines 6-26 and lines 60-62. Column 3, lines 1-3 recites, "the composition comprises a bioactive agent in combination with a plurality of polymers..." and column 5, lines 45-50, recites "The coating composition can also be used to coat stents...." Further, column 5, lines 58-62 states a range of about 0.05 mg to about 10 mg of bioactive agent per cm^2 of the gross surface of the device.

Given a stent range of 15-18 mm, this equates to $(0.05 \text{ mg/cm}^2) * (15-18 \text{ mm or } 1.5-1.8 \text{ cm}) = 0.075 \text{ mg to } 0.090 \text{ mg}$ This equals 75 ug to 90 ug, which falls within the range of 35-435 ug, as claimed.

However, Chudzik et al. do not explicitly recite, a compound having antiproliferative and anti-inflammatory properties. On the other hand, Morris et al.
teach the use of rapamycin which is known for its anti-proliferative and antiinflammatory properties. Therefore, it would be obvious to one with ordinary skill
in art to have the bioactive agent/compound of Chudzik et al. be rapamycin for
the purpose of its superior properties as an anti-proliferative and antiinflammatory for treating vascular remodeling. Further, this would fall within the
scope of bioactive agents described in column 5, lines 7-26 of Chudzik et al.

Regarding claims 2 and 9, further utilizing the compound to block proliferation of fibroblasts in a vascular wall in response to injury, thereby reducing a formation

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of vascular scar tissue would necessarily occur as a result of coating a stent with rapamycin in the dosages as described by Chudzik et al. with rapamycin as taught by Morris et al.

Regarding claims 3 and 10, Morris et al. teach rapamycin as discussed above with regard to claim 1.

Regarding claims 4 and 11, a compound that is analogs and cogeners that bind a high-affinity cytosolic protein, FKBP12, and possess pharmacologic properties equivalent to rapamycin, is also taught by Morris et al. as discussed with regard to claim 1 above.

Regarding claims 5 and 12, further utilizing the compound to affect a translation of certain proteins involved in a collagen formation or metabolism would necessarily occur as a result of coating a stent with rapamycin in the dosages as described by Chudzik et al. with rapamycin as taught by Morris et al.

Regarding claims 6 and 13, Morris et al. teach rapamycin as discussed above with regard to claim 1.

Regarding claims 7 and 17, a compound that is analogs and cogeners that bind a high-affinity cytosolic protein, FKBP12, and possess pharmacologic properties

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equivalent to rapamycin, is also taught by Morris et al. as discussed with regard to claim 1 above.

Double Patenting

4. The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. See *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970);and, *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 CFR 1.130(b).

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

5. Claims 1-14 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-11 of copending Application No. 09/850,233. Although the conflicting claims are not identical, they are not patentably distinct from each other because they are merely reworded representations for the same subject matter, perhaps slightly broader in some aspects while slightly more narrow in others.

This is a <u>provisional</u> obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

6. Claims 1-14 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-15 of copending Application No. 09/850,507. Although the conflicting claims are not identical,

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they are not patentably distinct from each other because they are merely reworded representations for the same subject matter, perhaps slightly broader in some aspects while slightly more narrow in others.

This is a <u>provisional</u> obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

7. Claims 1-14 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-17 of copending Application No. 09/850,232. Although the conflicting claims are not identical, they are not patentably distinct from each other because they are merely reworded representations for the same subject matter, perhaps slightly broader in some aspects while slightly more narrow in others.

This is a <u>provisional</u> obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

8. Claims 1-14 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-14 of copending Application No. 09/850,365. Although the conflicting claims are not identical, they are not patentably distinct from each other because they are merely reworded representations for the same subject matter, perhaps slightly broader in some aspects while slightly more narrow in others.

This is a <u>provisional</u> obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

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9. Claims 1-14 are provisionally rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-15 of copending Application No. 09/575,480. Although the conflicting claims are not identical, they are not patentably distinct from each other because they are merely reworded representations for the same subject matter, perhaps slightly broader in some aspects while slightly more narrow in others.

This is a <u>provisional</u> obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

10. Claims 1-14 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-20 of U.S. Patent No. 6,585,764. Although the conflicting claims are not identical, they are not patentably distinct from each other because they are merely reworded representations for the same subject matter, perhaps slightly broader in some aspects while slightly more narrow in others.

Conclusion

11. Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not

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mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Kathryn Odland whose telephone number is (703) 306-3454. The examiner can normally be reached on M-F (7:30-5:00) First Friday Off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Henry A Bennett can be reached on (703) 308-0101. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-

1113.

KO

Supervisory fatent Examiner